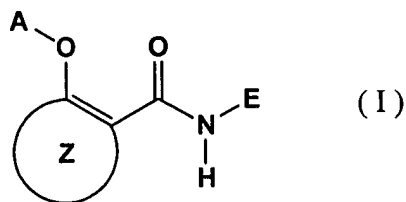


Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (Currently Amended): A method for ~~prophylactic or~~ therapeutic treatment of ~~tumor~~ skin cancer, melanoma, lung cancer, liver cancer, breast cancer, pancreatic cancer, leukemia, T-cell leukemia, rhabdomyoma, fibrosarcoma, or neuroblastoma in a mammal ~~including a human~~, which comprises the step of administering to a mammal a ~~prophylactically or therapeutically effective amount of a substance selected from the group consisting of~~ a compound represented by the following general formula (I) ~~or [[,]] a pharmacologically acceptable salt thereof, a hydrate thereof, and a solvate thereof:~~



wherein A represents a hydrogen atom or an acetyl group,

E represents a 2,5- or a 3,5-di-substituted phenyl group, wherein at least one of said substituents is a trifluoromethyl group, and the other substituent is selected from ~~the group consisting of~~

- a halogen atom,
- a nitro group,
- an alkyl group,
- a halogenated alkyl group,

an alkoxy group,
a halogenated alkoxy group,
an aryl-oxy group which may be substituted with one or more
substituents independently selected from ~~the group consisting of~~

a halogen atom,
an alkoxy group,
an alkyl group and
a cyano group,

an alkyl-sulfanyl group,
an alkoxy-carbonyl group,
a carboxy group, and

a monocyclic non-aromatic heterocyclic group which may be substituted with
one or more halogenated alkyl groups, or a mono-substituted thiazol-2-yl group;
or a di-substituted thiazol-2-yl group, wherein said substituents are
independently selected from ~~the group consisting of~~

a halogen atom,
an alkyl group which may be substituted with one or more substituents
independently selected from ~~the group consisting of~~
a carboxy group, and
an alkoxy-carbonyl group,
a halogenated alkyl group,
a cyano group,

an aryl group which may be substituted with one or more substituents independently selected from ~~the group consisting of~~

a halogen atom,

a halogenated alkyl group, and

an alkoxy group,

an alkyl-carbonyl group,

an alkoxy-carbonyl group,

a monocyclic non-aromatic heterocyclic group which may be substituted with one or more substituents independently selected from ~~the group consisting of~~

an alkyl group, and

an aryl group,

an aralkyl group,

an aryl-carbonyl group,

a carbamoyl group which may be substituted with one or more substituents independently selected from ~~the group consisting of~~

an alkyl group, and

an aralkyl group, and

a carboxy group,

ring Z represents a benzene ring which may have one or more substituents independently selected from ~~the group consisting of~~

a halogen atom,

a nitro group,

a cyano group,

an alkoxy group,

an alkyl group which may be substituted with one or more substituents

independently selected from ~~the group consisting of~~

a hydroxy group,

an aralkyl-oxy-imino group, and

an alkoxy-imino group,

an alkenyl group which may be substituted with one or more substituents

independently selected from ~~the group consisting of~~

an aryl group,

a cyano group,

an alkoxy-carbonyl group, and

a carboxy group,

an alkynyl group which may be substituted with one or more substituents

independently selected from ~~the group consisting of~~

an aryl group, and

a tri(alkyl)silyl group,

a halogenated alkyl group,

an aryl group which may be substituted with one or more substituents

independently selected from ~~the group consisting of~~

a halogen atom, and

a halogenated alkyl group,

an aralkyl group,

a monocyclic or a fused polycyclic heteroaryl group which may be substituted with one or more alkyl groups,

an alkyl-carbonyl group,

a monocyclic non-aromatic heterocyclic-carbonyl group which may be substituted with one or more aralkyl groups,

a monocyclic heteroaryl-sulfonyl group,

a carboxy group,

an alkoxy-carbonyl group,

a carbamoyl group which may be substituted with one or more substituents

independently selected from ~~the group consisting of~~

an aryl group which may be substituted with one or more halogenated

alkyl groups, and

an alkyl group,

a sulfamoyl group which may be substituted with one or more substituents

independently selected from ~~the group consisting of~~

an aryl group which may be substituted with one or more halogenated

alkyl groups, and

an alkyl group,

an amino group which may be substituted with one or more substituents

independently selected from ~~the group consisting of~~

an alkyl group,

an alkyl-carbonyl group,

an aryl-carbonyl group,

an alkyl-sulfonyl group, and
an aryl-sulfonyl group,
an ureido group which may be substituted with one or more aryl groups,
a thioureido group which may be substituted with one or more aryl groups,
a diazenyl group which may be substituted with one or more aryl groups wherein
said aryl groups may be substituted with one or more substituents independently
selected from ~~the group consisting of~~
a nitro group, and
a monocyclic heteroaryl-sulfamoyl group, and
a hydroxy group,
in addition to the group represented by formula –O-A and the group represented by
formula –CONH-E to a mammal.

2-28. (Canceled)

29. (Previously Presented) The method according to claim 1, wherein the mammal
is a human.

30-33. (Canceled)

34. (Currently Amended) A method for inhibiting proliferation of tumor cell or
cancer cell, which comprises ~~the step of~~ allowing an effective amount of a compound

according to claim 1 or a pharmacologically acceptable salt thereof to act on the tumor cell or cancer cell.

35. (Currently Amended) The method according to claim 1, wherein E is a 2,5- or 3,5-di-substituted phenyl group wherein at least one of said substituents is a trifluoromethyl group, and the other substituent is selected from ~~the group consisting of~~

- a halogen atom,
- a nitro group,
- an alkyl group,
- a halogenated alkyl group,
- an alkoxy group,
- a halogenated alkoxy group,
- an aryl-oxy group wherein said aryl-oxy group is a phenyl-oxy group or a naphthyl-oxy group, and said aryl-oxy group may be substituted with one or more substituents independently selected from ~~the group consisting of~~

- a halogen atom,
- an alkoxy group,
- an alkyl group and
- a cyano group,
- an alkyl-sulfanyl group,
- an alkoxy-carbonyl group,
- a carboxy group, and

a monocyclic non-aromatic heterocyclic group wherein said monocyclic non-aromatic heterocyclic group is a 1-pyrrolidinyl group, a piperidino group or a morpholino group, and said monocyclic non-aromatic heterocyclic group may be substituted with one or more halogenated alkyl group, or

a mono- or di-substituted thiazol-2-yl group wherein said substituents are independently selected from ~~the group consisting of~~

a halogen atom,

an alkyl group which may be substituted with one or more substituents independently selected from ~~the group consisting of~~

a carboxy group, and

an alkoxy-carbonyl group,

a halogenated alkyl group,

a cyano group,

a phenyl group which may be substituted with one or more substituents independently selected from ~~the group consisting of~~

a halogen atom,

a halogenated alkyl group, and

an alkoxy group,

an alkyl-carbonyl group,

an alkoxy-carbonyl group,

a monocyclic non-aromatic heterocyclic group wherein said monocyclic non-aromatic heterocyclic group is a piperidino group, a morpholino

group or a 1-piperazinyl group, and said monocyclic non-aromatic heterocyclic group may be substituted with one or more substituents independently selected from ~~the group consisting of~~

an alkyl group, and

a phenyl group,

a benzyl group,

a phenyl-carbonyl group,

a carbamoyl group which may be substituted with one or more substituents independently selected from ~~the group consisting of~~

an alkyl group, and

a phenyl-alkyl group, and

a carboxy group.

36. (Currently Amended) The method according to claim 1, wherein E [[a]] is a 3,5-bis(trifluoromethyl)phenyl group, a 3-fluoro-5-(trifluoromethyl)phenyl group, a 3-bromo-5-(trifluoromethyl)phenyl group, a 3-methoxy-5-(trifluoromethyl)phenyl group, a 3-methoxycarbonyl-5-(trifluoromethyl)phenyl group, a 3-carboxy-5-(trifluoromethyl)phenyl group, a 2-chloro-5-(trifluoromethyl)phenyl group, a 2,5-bis(trifluoromethyl)phenyl group, a 2-fluoro-5-(trifluoromethyl)phenyl group, a 2-nitro-5-(trifluoromethyl)phenyl group, a 2-methyl-5-(trifluoromethyl)phenyl group, a 2-methoxy-5-(trifluoromethyl)phenyl group, a 2-methylsulfanyl-5-(trifluoromethyl)phenyl group, a 2-(1-pyrrolidinyl)-5-(trifluoromethyl)phenyl group, a 2-morpholino-5-(trifluoromethyl)phenyl group, a 2-bromo-5-(trifluoromethyl)phenyl group, a 2-(2-

naphthyloxy)-5-(trifluoromethyl)phenyl group, a 2-(2,4-dichlorophenoxy)-5-(trifluoromethyl)phenyl group, a 2-[4-(trifluoromethyl)piperidin-1-yl]-5-(trifluoromethyl)phenyl group, a 2-(2,2,2-trifluoroethoxy)-5-(trifluoromethyl)phenyl group, a 2-(2-methoxyphenoxy)-5-(trifluoromethyl)phenyl group, a 2-(4-chloro-3,5-dimethylphenoxy)-5-(trifluoromethyl)phenyl group, a 2-piperidino-5-(trifluoromethyl)phenyl group, a 2-(4-methylphenoxy)-5-(trifluoromethyl)phenyl group, a 2-(4-chlorophenoxy)-5-(trifluoromethyl)phenyl group, a 2-(4-cyanophenoxy)-5-(trifluoromethyl)phenyl group, a 2-(4-methoxyphenoxy)-5-(trifluoromethyl)phenyl group, a 5-bromo-4-[(1,1-dimethyl)ethyl]thiazol-2-yl group, a 5-bromo-4-(trifluoromethyl)thiazol-2-yl group, a 5-cyano-4-[(1,1-dimethyl)ethyl]thiazol-2-yl group, a 5-methylthiazol-2-yl group, a 4,5-dimethylthiazol-2-yl group, a 5-methyl-4-phenylthiazol-2-yl group, a 5-(4-fluorophenyl)-4-methylthiazol-2-yl group, a 4-methyl-5-[3-(trifluoromethyl)phenyl]thiazol-2-yl group, a 4-[(1,1-dimethyl)ethyl]-5-ethylthiazol-2-yl group, a 4-ethyl-5-phenylthiazol-2-yl group, a 4-isopropyl-5-phenylthiazol-2-yl group, a 4-butyl-5-phenylthiazol-2-yl group, a 4-[(1,1-dimethyl)ethyl]-5-[(2,2-dimethyl)propionyl]thiazol-2-yl group, a 4-[(1,1-dimethyl)ethyl]-5-(ethoxycarbonyl)thiazol-2-yl group, a 4-[(1,1-dimethyl)ethyl]-5-piperidinethiazol-2-yl group, a 4-[(1,1-dimethyl)ethyl]-5-morpholinethiazol-2-yl group, a 4-[(1,1-dimethyl)ethyl]-5-(4-methylpiperazin-1-yl)thiazol-2-yl group, a 4-[(1,1-dimethyl)ethyl]-5-(4-phenylpiperazin-1-yl)thiazol-2-yl group, a 5-carboxymethyl-4-phenylthiazol-2-yl group, a 4,5-diphenylthiazol-2-yl group, a 4-benzyl-5-phenylthiazol-2-yl group, a 5-phenyl-4-(trifluoromethyl)thiazol-2-yl group, a 5-acetyl-4-phenylthiazol-2-yl group, a 5-benzoyl-4-phenylthiazol-2-yl group, a 5-ethoxycarbonyl-4-phenylthiazol-2-yl group, a 5-

ethoxycarbonyl-4-(pentafluorophenyl)thiazol-2-yl group, a 5-methylcarbamoyl-4-phenylthiazol-2-yl group, a 5-ethylcarbamoyl-4-phenylthiazol-2-yl group, a 5-isopropylcarbamoyl-4-phenylthiazol-2-yl group, a 5-(2-phenylethyl)carbamoyl-4-phenylthiazol-2-yl group, a 5-ethoxycarbonyl-4-(trifluoromethyl)thiazol-2-yl group, a 5-carboxy-4-[(1,1-dimethyl)ethyl]thiazol-2-yl group, a 5-(ethoxycarbonyl)methyl-4-phenylthiazol-2-yl group, a 5-carboxy-4-phenylthiazol-2-yl group, a 5-propylcarbamoyl-4-phenylthiazol-2-yl group, a 4-[(1,1-dimethyl)ethyl]thiazol-2-yl group, a 4-phenylthiazol-2-yl group, a 4-[3,5-bis(trifluoromethyl)phenyl]thiazol-2-yl group, a 4-(2,4-dichlorophenyl)thiazol-2-yl group, a 4-(3,4-dichlorophenyl)thiazol-2-yl group, a 4-[4-(trifluoromethyl)phenyl]thiazol-2-yl group, a 4-(2,5-difluorophenyl)thiazol-2-yl group, a 4-(4-methoxyphenyl)thiazol-2-yl group, a 4-[3-(trifluoromethyl)phenyl]thiazol-2-yl group, a or 4-(pentafluorophenyl)thiazol-2-yl group.

37. (Currently Amended) The method according to claim 1, wherein ring Z is a benzene ring which may have one or more substituents independently selected from ~~the~~ group consisting of

a halogen atom,

a nitro group,

a cyano group,

an alkoxy group,

an alkyl group which may be substituted with one or more substituents

independently selected from ~~the group consisting of~~

a hydroxyl group,

an benzyl-oxy-imino group, and

an alkoxy-imino group,

an alkenyl group which may be substituted with one or more substituents independently selected from ~~the group consisting of~~

a phenyl group,

a cyano group,

an alkoxy-carbonyl group, and

a carboxy group,

an alkynyl group which may be substituted with one or more substituents independently selected from ~~the group consisting of~~

a phenyl group, and

a tri(alkyl)silyl group,

a halogenated alkyl group,

a phenyl group which may be substituted with one or more substituents independently selected from ~~the group consisting of~~

a halogen atom, and

a halogenated alkyl group,

a phenyl-alkyl group,

a monocyclic or a fused polycyclic heteroaryl group wherein said monocyclic or a fused polycyclic heteroaryl group is a 1-pyrrolyl group, a 2-pyrrolyl group, a 3-pyrrolyl group, a 2-thienyl group, a 3-thienyl group, a 2-thiazolyl group, a 4-thiazolyl group, a 5-thiazolyl group, a 2-pyridyl group, a 3-pyridyl group, a 4-pyridyl group or an imidazo[1,2-a]pyridin-2-yl group, and said monocyclic or a

fused polycyclic heteroaryl group may be substituted with one or more alkyl groups,

an alkyl-carbonyl group,

a monocyclic non-aromatic heterocyclic-carbonyl group wherein said monocyclic non-aromatic heterocyclic-carbonyl group is a piperidino-carbonyl group, and said monocyclic non-aromatic heterocyclic-carbonyl group may be substituted with one or more benzyl groups,

a monocyclic heteroaryl-sulfonyl group wherein said monocyclic heteroaryl-sulfonyl group is a 1-pyrrolyl-sulfonyl group, a 2-pyrrolyl-sulfonyl group or a 3-pyrrolyl-sulfonyl group,

a carboxy group,

an alkoxy-carbonyl group,

a carbamoyl group which may be substituted with one or more substituents independently selected from ~~the group consisting of~~

- a phenyl group which may be substituted with one or more halogenated alkyl groups, and
- an alkyl group,

a sulfamoyl group which may be substituted with one or more substituents independently selected from ~~the group consisting of~~

- a phenyl group which may be substituted with one or more halogenated alkyl groups, and
- an alkyl group,

an amino group which may be substituted with one or more substituents independently selected from ~~the group consisting of~~

- an alkyl group,
- an alkyl-carbonyl group,
- a phenyl-carbonyl group,
- an alkyl-sulfonyl group, and
- a phenyl-sulfonyl group,

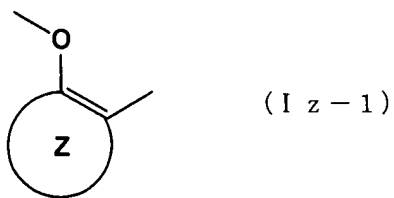
an ureido group which may be substituted with one or more phenyl groups,
a thioureido group which may be substituted with one or more phenyl groups,
a diazenyl group which may be substituted with one or more phenyl groups
wherein said phenyl groups may be substituted with one or more substituents independently selected from ~~the group consisting of~~

- a nitro group, and
- a monocyclic heteroaryl-sulfamoyl group wherein said monocyclic heteroaryl-sulfamoyl group is a 2-pyridyl-sulfamoyl group, a 3-pyridyl-sulfamoyl group or a 4-pyridyl-sulfamoyl group, and

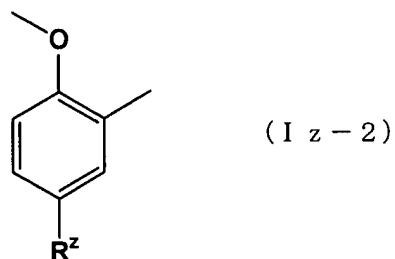
a hydroxy group,

in addition to the group represented by formula —O—A wherein A has the same meaning as that defined above and the group represented by formula —X—E wherein each of X and E has the same meaning as that defined above.

38. (Currently Amended) The method according to claim 1, wherein the following partial formula (Iz-1) in the general formula containing ring Z



is represented by the following formula (Iz-2):



wherein R^Z represents

a hydrogen atom,

a halogen atom,

a nitro group,

a cyano group,

an alkoxy group,

an alkyl group which may be substituted with one or more substituents

independently selected from ~~the group consisting of~~

a hydroxy group,

an aralkyl-oxy-imino group, and

an alkoxy-imino group,

an alkenyl group which may be substituted with one or more substituents

independently selected from ~~the group consisting of~~

an aryl group,

a cyano group,

an alkoxy-carbonyl group, and

a carboxy group,

an alkynyl group which may be substituted with one or more substituents independently selected from ~~the group consisting of~~

an aryl group, and

a tri(alkyl)silyl group,

a halogenated alkyl group,

an aryl group which may be substituted with one or more substituents independently selected from ~~the group consisting of~~

a halogen atom, and

a halogenated alkyl group,

an aralkyl group,

a monocyclic or a fused polycyclic heteroaryl group which may be substituted with one or more alkyl groups,

an alkyl-carbonyl group,

a monocyclic non-aromatic heterocyclic-carbonyl group which may be substituted with one or more aralkyl groups,

a monocyclic heteroaryl-sulfonyl group,

a carboxy group,

an alkoxy-carbonyl group,

a carbamoyl group which may be substituted with one or more substituents independently selected from ~~the group consisting of~~

an aryl group which may be substituted with one or more halogenated alkyl groups, and
an alkyl group,
a sulfamoyl group which may be substituted with one or more substituents independently selected from ~~the group consisting of~~
an aryl group which may be substituted with one or more halogenated alkyl groups, and
an alkyl group,
an amino group which may be substituted with one or more substituents independently selected from ~~the group consisting of~~
an alkyl group,
an alkyl-carbonyl group,
an aryl-carbonyl group,
an alkyl-sulfonyl group, and
an aryl-sulfonyl group,
an ureido group which may be substituted with one or more aryl groups,
a thioureido group which may be substituted with one or more aryl groups, or
a diazenyl group which may be substituted with one or more aryl groups wherein said aryl groups may be substituted with one or more substituents independently selected from ~~the group consisting of~~
a nitro group, and
a monocyclic heteroaryl-sulfamoyl group.

39. (Currently Amended) The use according to claim 38, wherein R^z is

a hydrogen atom,

a halogen atom,

a nitro group,

a cyano group,

an alkoxy group,

an alkyl group which may be substituted with one or more substituents

independently selected from ~~the group consisting of~~

a hydroxyl group,

an benzyl-oxy-imino group, and

an alkoxy-imino group,

an alkenyl group which may be substituted with one or more substituents

independently selected from ~~the group consisting of~~

a phenyl group,

a cyano group,

an alkoxy-carbonyl group, and

a carboxy group,

an alkynyl group which may be substituted with one or more substituents

independently selected from ~~the group consisting of~~

a phenyl group, and

a tri(alkyl)silyl group,

a halogenated alkyl group,

a phenyl group which may be substituted with one or more substituents

independently selected from ~~the group consisting of~~

a halogen atom, and

a halogenated alkyl group,

a phenyl-alkyl group,

a monocyclic or a fused polycyclic heteroaryl group wherein said monocyclic or

a fused polycyclic heteroaryl group is a 1-pyrrolyl group, a 2-pyrrolyl group, a 3-

pyrrolyl group, a 2-thienyl group, a 3-thienyl group, a 2-thiazolyl group, a 4-

thiazolyl group, a 5-thiazolyl group, a 2-pyridyl group, a 3-pyridyl group, a 4-

pyridyl group or an imidazo[1,2-a]pyridin-2-yl group, and said monocyclic or a

fused polycyclic heteroaryl group may be substituted with one or more alkyl

groups,

an alkyl-carbonyl group,

a monocyclic non-aromatic heterocyclic-carbonyl group wherein said

monocyclic non-aromatic heterocyclic-carbonyl group is a piperidino-carbonyl

group, and said monocyclic non-aromatic heterocyclic-carbonyl group may be

substituted with one or more benzyl groups,

a monocyclic heteroaryl-sulfonyl group wherein said monocyclic heteroaryl-

sulfonyl group is a 1-pyrrolyl-sulfonyl group, a 2-pyrrolyl-sulfonyl group or a 3-

pyrrolyl-sulfonyl group,

a carboxy group,

an alkoxy-carbonyl group,

a carbamoyl group which may be substituted with one or more substituents independently selected from ~~the group consisting of~~

a phenyl group which may be substituted with one or more halogenated alkyl groups, and
an alkyl group,

a sulfamoyl group which may be substituted with one or more substituents independently selected from ~~the group consisting of~~

a phenyl group which may be substituted with one or more halogenated alkyl groups, and
an alkyl group,

an amino group which may be substituted with one or more substituents independently selected from ~~the group consisting of~~

an alkyl group,
an alkyl-carbonyl group,
a phenyl-carbonyl group,
an alkyl-sulfonyl group, and
a phenyl-sulfonyl group,

an ureido group which may be substituted with one or more phenyl groups,
a thioureido group which may be substituted with one or more phenyl groups, or
a diazenyl group which may be substituted with one or more phenyl groups wherein said phenyl groups may be substituted with one or more substituents independently selected from ~~the group consisting of~~

a nitro group, and

a monocyclic heteroaryl-sulfamoyl group wherein said monocyclic heteroaryl-sulfamoyl group is a 2-pyridyl-sulfamoyl group, a 3-pyridyl-sulfamoyl group or a 4-pyridyl-sulfamoyl group.

40. (Currently Amended) The method according to claim 38, wherein R^z is a hydrogen atom, a halogen atom, a nitro group, a cyano group, a methoxy group, a methyl group, an isopropyl group, a tert-butyl group, a 1,1,3,3-tetramethylbutyl group, a 2-phenylethen-1-yl group, a 2,2-dicyanoethen-1-yl group, a 2-cyano-2-(methoxycarbonyl)ethen-1-yl group, a 2-carboxy-2-cyanoethen-1-yl group, an ethynyl group, a phenylethynyl group, a (trimethylsilyl)ethynyl group, a trifluoromethyl group, a pentafluoroethyl group, a phenyl group, a 4-(trifluoromethyl)phenyl group, a 4-fluorophenyl group, a 2,4-difluorophenyl group, a 2-phenethyl group, a 1-hydroxyethyl group, a 1-(methoxyimino)ethyl group, a 1-[(benzyloxy)imino]ethyl group, a 2-thienyl group, a 3-thienyl group, a 1-pyrrolyl group, a 2-methylthiazol-4-yl group, an imidazo[1,2-a]pyridin-2-yl group, a 2-pyridyl group, [[a]] an acetyl group, an isobutyryl group, a piperidinocarbonyl group, a 4-benzylpiperidinocarbonyl group, a (pyrrol-1-yl)sulfonyl group, a carboxy group, a methoxycarbonyl group, an N-[3,5-bis(trifluoromethyl)phenyl]carbamoyl group, an N,N-dimethylcarbamoyl group, a sulfamoyl group, an N-[3,5-bis(trifluoromethyl)phenyl]sulfamoyl group, an N,N-dimethylsulfamoyl group, an amino group, an N,N-dimethylamino group, an acetylamino group, a benzoylamino group, a methanesulfonylamino group, a benzenesulfonylamino group, a 3-phenylureido group, a (3-phenyl)thioureido group, a (4-nitrophenyl)diazenyl group, [[a]] or a {[4-(pyridin-2-yl)sulfamoyl]phenyl} diazenyl group.

41. (Currently Amended) The method according to claim 1, wherein

A is hydrogen atom,

E is a 2,5- or 3,5-di-substituted phenyl group wherein at least one of said substituents is a trifluoromethyl group, and the other substituent is selected from ~~the group consisting of~~

a halogen atom,

a halogenated alkyl group,

an alkoxy group, or

a di-substituted thiazol-2-yl group wherein said substituents are independently selected from ~~the group consisting of~~

an alkyl group,

a halogenated alkyl group,

a cyano group,

an aryl group,

an alkyl-carbonyl group,

a monocyclic non-aromatic heterocyclic group which may be substituted with an aryl group, and

an aryl-carbonyl group,

ring Z is a benzene ring which may have one or more substituents independently selected from ~~the group consisting of~~

a halogen atom,

an alkyl group,

an alkenyl group which may be substituted with an aryl group,
a halogenated alkyl group,
an aryl group, and
a monocyclic heteroaryl group,
in addition to the group represented by formula —O—A and the group represented by
formula —CONH—E .